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(71) Applicants (for all designated States except US): **TSINGHUA UNIVERSITY** [CN/CN]; Tsinghua University, Haidian District, Beijing 100084 (CN). **CAPITAL BIOCHIP COMPANY, LTD.** [CN/CN]; Jia N° 2, Qinghua West Road, Haidian District, Beijing 100084 (CN).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **YI, Guangshun** [CN/CN]; Tsinghua University, Haidian District, Beijing 100084 (CN). **SUN, Baoquan** [CN/CN]; Tsinghua University, Haidian District, Beijing 100084 (CN). **CHEN, Depu** [CN/CN]; Tsinghua University, Haidian District, Beijing 100084 (CN). **ZHOU, Yuxiang** [CN/CN]; Tsinghua University, Haidian District, Beijing 100084 (CN). **CHENG, Jing** [CN/CN]; Tsinghua University, Haidian District, Beijing 100084 (CN). **YANG, Wenjun** [CN/CN]; Tsinghua University, Haidian District, Beijing

100084 (CN). **GE, Yue** [CN/CN]; Tsinghua University, Haidian District, Beijing 100084 (CN). **GUO, Lianghong** [CN/CN]; Tsinghua University, Haidian District, Beijing 100084 (CN).

(74) Agent: **JEEKAI & PARTNERS**; Suite 602, Jinyu Tower, A129 West Xuan Wu Men Street, Xicheng District, Beijing 100031 (CN).

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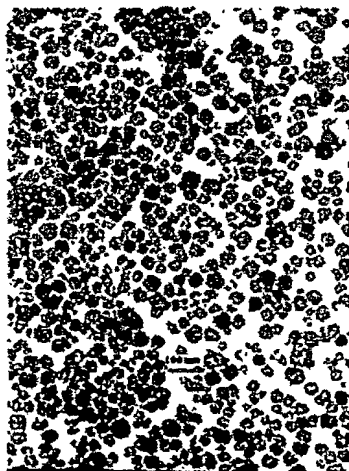
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(54) Title: NANOMETER-SIZED UP-CONVERTING PHOSPHOR FLUORIDE PARTICLES AND PROCESS OF PREPARATION



(57) Abstract: Nanometer-scaled up-converting fluoride phosphor particles and processes of making them are disclosed. In the process, an aqueous solution consisting of soluble salts of rare-earth metal ions at a molar ratio of (yttrium, lanthanum or gadolinium): (ytterbium: (erbium, holmium, terbium or thulium) = (70-90):(0-29):(0.001-15) is mixed a rare-earth metal chelator and a soluble fluoride salt to form precipitates, which are then annealed at an elevated temperature to produce nanometer-scaled up-converting fluoride phosphor particles. The particle size is between 35nm and 200nm, and can be controlled by the amount of the metal chelator added to the solution. The nanometer-sized particle is applicable to many biological assays.

NANOMETER-SIZED UP-CONVERTING PHOSPHOR FLUORIDE PARTICLES
AND PROCESS OF PREPARATION

Technical Field

The present invention relates to nanometer-sized fluorescent particles and process of making them. More particularly, the present invention relates to rare-earth ion doped, nanometer-sized up-converting phosphor fluoride particles and process of making them.

Background Art

Up-converting phosphor is a type of material that emits visible light when excited by long wavelength light. Ytterbium/erbium co-doped fluoride is a high efficiency up-converting phosphor that emits strong green fluorescence and relatively weak red fluorescence when excited by 946-970nm infrared light.

Due to their unique properties, up-converting phosphors are used in the fabrication of light-emitting diodes, solid-state lasers, and as ultra-sensitive fluorescent labels in biological detections (U.S. Patent No. 5,674,698).

As a fluorescent label for biological molecules, the phosphor must be small in size and uniform, highly luminescent, and stable in aqueous environment. We have disclosed a method of preparing nanometer-sized molybdenum up-converting phosphor particles (Patent Number: CN 01134861.5). With this method, particles of 50-60nm in diameter were prepared. They were easy to make, and highly luminescent. But the range of particle size was limited to 50-60nm. And the annealing temperature is high. James Kane has disclosed in US Patent 5,891,361 and WO 97/46488 a process for preparing small up-converting fluoride phosphor particles. The process involves high temperature (750°C) heating, and the size of the particle (about 400nm) thus obtained is too large for many biological applications.

Disclosure of the Invention

The present invention relates to nanometer-sized fluorescent particles and process of making them.

The aim of this invention is to disclose nanometer-scaled up-converting fluoride phosphor particles and processes of making them. The size of the phosphor particles can be varied over a range of about 35nm to 350nm in a controlled fashion, which is significantly smaller than those disclosed in prior arts. The particles are uniform in size, and highly

luminescent. In addition, the phosphor particles are prepared at annealing temperatures much lower than those employed in prior arts. Therefore, the nanometer-scaled up-converting fluoride phosphor particles disclosed in this application are more applicable to biological detections.

5 In one aspect, the present invention is directed to a phosphor fluoride particle that emits light in the visible wavelength range when excited by long wavelength light that has a uniform particle size of less than 350 nanometers.

In another aspect, the present invention is directed to a process of preparing a phosphor fluoride particle that emits light in the visible wavelength range when excited by
10 long wavelength light that has a uniform particle size of less than 350 nanometers, which process comprises: a) preparing an aqueous solution of soluble salts of a phosphor host, an absorber/emitter pair and a rare-earth metal chelator; and b) contacting said prepared aqueous solution of soluble salts of said phosphor host, absorber/emitter pair and rare-earth metal chelator with an aqueous fluoride-containing compound at a temperature ranging from
15 about 0°C to about 100°C for a sufficient time to obtain a precipitate of a phosphor fluoride particle; and c) heating said precipitate at a temperature ranging from about 300°C to about 450°C for a time ranging from about 1 hour to about 10 hours to obtain a phosphor fluoride particle that emits light in the visible wavelength range when excited by long wavelength light that has a uniform particle size of less than 350 nanometers.

20 **Brief Description of the Drawings**

Figure 1 illustrates a transmission electron micrograph (TEM) of phosphor fluoride particles with a uniform particle size of about 37 nanometers prepared by the process described in Example 1.

Figure 2 illustrates a TEM of phosphor fluoride particles with a uniform particle size
25 of about 43 nanometers prepared by the process described in Example 2.

Figure 3 illustrates a TEM of phosphor fluoride particles with a uniform particle size of about 77 nanometers prepared by the process described in Example 3.

Figure 4 illustrates a TEM of phosphor fluoride particles with a uniform particle size of about 96 nanometers prepared by the process described in Example 4.

Figure 5 illustrates a TEM of phosphor fluoride particles with a uniform particle size of about 166 nanometers prepared by the process described in Example 5.

Figure 6 illustrates up-converting fluorescent spectrum of phosphor fluoride particles with a uniform particle size of about 37 nanometers prepared by the process described in

5 Example 1.

Modes of Carrying Out the Invention

For clarity of disclosure, and not by way of limitation, the detailed description of the invention is divided into the subsections that follow.

A. Definitions

10 Unless defined otherwise, all technical and scientific terms used herein have the same meaning as is commonly understood by one of ordinary skill in the art to which this invention belongs. All patents, applications, published applications and other publications referred to herein are incorporated by reference in their entirety. If a definition set forth in this section is contrary to or otherwise inconsistent with a definition set forth in the patents,
15 applications, published applications and other publications that are herein incorporated by reference, the definition set forth in this section prevails over the definition that is incorporated herein by reference.

As used herein, "a" or "an" means "at least one" or "one or more."

As used herein, "a uniform particle size" means that a group of particles has a
20 standard deviation of particle sizes that equals to or is less than 20%. Preferably, the group of particles has a standard deviation of particle sizes that equals to or is less than 15%, 10%, 5%, 2%, 1%, or less than 1%.

As used herein, "phosphor" means solid, inorganic, crystalline material that shows luminescence upon optical excitation.

25 As used herein, "phosphor host" means one (and usually the major one) of the three components in a phosphor which does not participate in the light absorption or light emission process, but provides structural environment for the other two components.

As used herein, "absorber/emitter pair" means the two components in a phosphor which respectively absorbs longer-wavelength light and emits shorter-wavelength light to
30 complete the up-converting process.

B. Phosphor fluoride particles

In one aspect, the present invention is directed to a phosphor fluoride particle that emits light in the visible wavelength range when excited by long wavelength light that has a uniform particle size of less than 350 nanometers.

5 Preferably, the phosphor fluoride particle comprises a phosphor host and an absorber-emitter pair. Any suitable phosphor host, *e.g.*, yttrium, lanthanum or gadolinium, can be used in the present phosphor fluoride particle. Any suitable absorber, *e.g.*, ytterbium, can be used in the present phosphor fluoride particle. Any suitable emitter, *e.g.*, erbium, holmium, terbium or thulium, can be used in the present phosphor fluoride particle.

10 In a specific embodiment, the absorber is ytterbium and the emitter is erbium, holmium, terbium or thulium.

In another specific embodiment, the particle has a molar ratio:

(yttrium, lanthanum or gadolinium):ytterbium:(erbium, holmium, terbium or thulium)
= (70-90):(0-29):(0.001-15).

15 In still another specific embodiment, the phosphor fluoride particle has a formula of $\text{YF}_3\text{:Yb,Er}$.

In still another specific embodiment, the phosphor fluoride particle has a formula of $\text{NaYF}_4\text{:Yb,Er}$.

The phosphor fluoride particle can have any size that is less than 350 nanometers.

20 For example, the phosphor fluoride particle can have a particle size ranging from about 35 nanometers to about 200 nanometers.

In yet another specific embodiment, the phosphor fluoride particle can further comprise a transparent coating layer. The coated particle can further comprise a layer of immobilized biological moiety, *e.g.*, a biological molecule. Exemplary biological moieties

25 include a cell, a cellular organelle, a virus, a molecule and an aggregate or complex thereof.

The phosphor fluoride particle can have any suitable shape, *e.g.*, a rectangle, a circle, an ellipse, or other regular or irregular shapes. Preferably, phosphor fluoride particle has a spherical shape.

C. Processes of preparing phosphor fluoride particles

In another aspect, the present invention is directed to a process of preparing a phosphor fluoride particle that emits light in the visible wavelength range when excited by long wavelength light that has a uniform particle size of less than 350 nanometers, which process comprises: a) preparing an aqueous solution of soluble salts of a phosphor host, an absorber/emitter pair and a rare-earth metal chelator; and b) contacting said prepared aqueous solution of soluble salts of said phosphor host, absorber/emitter pair and rare-earth metal chelator with an aqueous fluoride-containing compound at a temperature ranging from about 0°C to about 100°C for a sufficient time to obtain a precipitate of a phosphor fluoride particle; and c) heating said precipitate at a temperature ranging from about 300°C to about 450°C for a time ranging from about 1 hour to about 10 hours to obtain a phosphor fluoride particle that emits light in the visible wavelength range when excited by long wavelength light that has a uniform particle size of less than 350 nanometers.

Any suitable phosphor host can be used in the present processes. For example, yttrium, lanthanum or gadolinium can be used as the phosphor host in the present processes. Any suitable absorber can be used in the present processes. For example, ytterbium can be used as the absorber in the present processes. Any suitable emitter can be used in the present processes. For example, erbium, holmium, terbium or thulium can be used as the emitter in the present processes. In a specific embodiment, the absorber is ytterbium and the emitter is erbium, holmium, terbium or thulium.

Any suitable rare-earth metal chelator can be used in the present processes. For example, ethylenediaminetetraacetic acid, triethylenetetraaminhexaacetic acid, diethylenetriaminepentaacetic acid, hydroxyethylethylenediaminetriacetic acid, 1,2-diaminocyclohexanetetraacetic acid, ethylene glycol bis (b-aminoethylether) tetraacetic acid and a salt thereof can be used as the rare-earth metal chelator in the present processes.

Any suitable aqueous fluoride-containing compound can be used in the present processes. For example, NaF, KF, NH₄F and HF can be used as the aqueous fluoride-containing compound in the present processes. The aqueous fluoride-containing compound can be contained in an aqueous solution prior to or concurrently with contacting with the prepared aqueous solution of soluble salts of the phosphor host, the absorber/emitter pair and the rare-earth metal chelator. The amount of the rare-earth metal chelator and the amount of

total rare-earth ions in the aqueous solution can have any suitable ratios. For example, the amount of the rare-earth metal chelator can be about 0-1 times the amount of total rare-earth ions in the aqueous solution.

The soluble salts of the phosphor host and the absorber/emitter pair can be prepared
5 by any suitable methods. For example, the soluble salts of the phosphor host and the absorber/emitter pair can be obtained by dissolving the corresponding metal oxide in hydrochloric acid or nitric acid and subsequently removing the residual acid.

The present process can further comprise coating the prepared phosphor fluoride particle with a transparent layer, *e.g.*, a SiO₂ layer. The present process can further comprise
10 immobilizing any desirable moieties, *e.g.*, biological molecules, to the phosphor fluoride particle. For example, the present process can further comprise immobilizing a biological molecule to the SiO₂-protected particle. Any suitable biological molecule can be used. Exemplary biological molecules include an amino acid, a peptide, a protein, a nucleoside, a nucleotide, an oligonucleotide, a nucleic acid, a vitamin, a monosaccharide, an
15 oligosaccharide, a carbohydrate, a lipid and a complex thereof.

In still another aspect, the present invention is directed to a phosphor fluoride particle that is prepared by the above-described process. Preferably, the phosphor fluoride particle has a molar ratio: (yttrium, lanthanum or gadolinium):ytterbium:(erbium, holmium, terbium or thulium) = (70-90):(0-29):(0.001-15). Preferably, the phosphor fluoride particle has a
20 formula of YF₃:Yb,Er. Also preferably, the phosphor fluoride particle has a formula of NaYF₄:Yb,Er. Still preferably, the phosphor fluoride particle has a particle size from about 35 nanometers to about 200 nanometers.

D. Examples

The nanometer-scaled up-converting fluoride phosphor particles prepared in the
25 process disclosed in the invention has the following characteristics: (a) The particle size is on the nanometer scale and as small as 37nm. (b) The particle size can be varied between 37nm and 166nm by controlling the amount of the rare-earth metal chelator. (c) The up-converting fluorescence of the fluoride phosphor is four times higher than that of the molybdenum phosphor. (d) The particles are prepared at a much lower annealing

temperature. (e) The process is highly reproducible. Variation of the particle size between two preparations is less than 5%.

Example 1: Preparation of 37nm Up-Converting Fluoride Phosphors

(1) Yttrium chloride (YCl_3) was obtained by dissolving 11.2905 g of yttrium oxide (Y_2O_3) in hydrochloric acid (HCl) or nitric acid (HNO_3). The mixture was heated and the solvent was let evaporated. Five hundred milliliters of de-ionized water was then added to re-dissolve the solid to yield a 0.2 M YCl_3 stock solution. Five hundred milliliters of 0.2 M stock solutions of gadolinium chloride (GdCl_3), lanthanum chloride (LaCl_3), ytterbium chloride (YbCl_3), erbium chloride (ErCl_3), thulium chloride (TmCl_3), and holmium chloride (HoCl_3) were prepared in the same way from 18.125 g of Gd_2O_3 , 16.3 g of La_2O_3 , 19.7 g of Yb_2O_3 , 19.1 g of Er_2O_3 , 19.3 g of Tm_2O_3 and 18.9 g of Ho_2O_3 , respectively. Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 6 mL: 3.4 mL: 0.6 mL and added into a 100 mL three-necked Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 80:17:3$.

(2) 20 mL of 0.2M ethylenediaminetetraacetic acid (EDTA) solution was introduced to the solution prepared in (1).

(3) A sodium fluoride (NaF) solution was prepared by dissolving 2.1 g of NaF in 60 mL of de-ionized water. The solution prepared in (2) was injected into the NaF solution. The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried.

(5) The precursors obtained in (4) were heated at 400°C for 4 h. The nanometer-sized up-converting fluorescent particles were obtained with an average diameter of 37 nm.

Example 2: Preparation of 43nm Up-Converting Fluoride Phosphors

(1) Stock solutions of YCl_3 , YbCl_3 and TmCl_3 were mixed at a volume ratio of 17.4 mL: 2.6 mL: 0.02 mL and added into a 100 mL three-necked Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 87:13:0.1$.

(2) 15 mL of 0.2 M EDTA solution was introduced to the solution prepared in (1).

(3) A NaF solution was prepared by dissolving 2.1 g of NaF in 65 mL of de-ionized water. The solution prepared in (2) was injected into the NaF solution. The mixture was stirred vigorously for 1 h at room temperature.

5 (4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried.

(5) The precursors obtained in (4) were heated at 400°C for 4 h. The nanometer-sized up-converting fluorescent particles were obtained with an average diameter of 43 nm.

Example 3: Preparation of 77nm Up-Converting Fluoride Phosphors

10 (1) Stock solutions of YCl_3 , YbCl_3 and HoCl_3 were mixed at a volume ratio of 16 mL: 3.86 mL: 0.04 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 80:19.8:0.2$.

(2) 10 mL of 0.2 M EDTA solution was introduced to the solution prepared in (1).

15 (3) A solution of NaF was prepared by dissolving 2.1 g of NaF in 70 mL of de-ionized water. The solution prepared in (2) was injected into the NaF solution. The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried.

(5) The precursors obtained in (4) were heated at 400°C for 5 h. The nanometer-sized up-converting fluorescent particles were obtained with an average diameter of 77 nm.

20 Example 4: Preparation of 96nm Up-Converting Fluoride Phosphors

(1) Stock solutions of GdCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 14 mL: 5.8 mL: 0.2 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 70:29:1$.

(2) 5 mL of 0.2 M EDTA solution was introduced to the solution prepared in (1).

25 (3) A solution of ammonium fluoride (NH_4F) was prepared by dissolving 2.1 g of NH_4F in 75 mL of de-ionized water. The solution prepared in (2) was injected into the NH_4F solution. The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried.

(5) The precursors obtained in (4) were heated at 350°C for 5 h. The nanometer-sized up-converting fluorescent particles were obtained with an average diameter of 96 nm.

Example 5: Preparation of 166nm Up-Converting Fluoride Phosphors

5 (1) Stock solutions of LaCl_3 and ErCl_3 were mixed at a volume ratio of 18 mL: 2 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{La}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 90:0:10$.

(2) A solution of potassium fluoride (KF) was prepared by dissolving 2.1 g of KF in 75 mL of de-ionized water. The solution prepared in (2) was injected into the KF solution. The mixture was stirred vigorously for 1 h at room temperature.

10 (3) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried.

(4) The precursors obtained in (4) were annealed heated at 400°C for 5 h. The nanometer-sized up-converting fluorescent particles were obtained with average diameter of 166 nm.

15 Example 6: Preparation of 43nm Up-Converting Fluoride Phosphor with TTHA Chelator

(1) Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 16 mL: 3.6 mL: 0.4 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 80:18:2$.

20 (2) 5 mL of 0.2 M triethylenetetraaminhexaacetic acid (TTHA) solution was introduced to the solution prepared in (1).

(3) A solution of NaF was prepared by dissolving 2.1 g of NaF in 75 mL of de-ionized water. The solution described in (2) was injected into the NaF solution at 50°C. The mixture was stirred vigorously for 1 h at room temperature.

25 (4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried. The average diameter of the precursor particles was 43 nm.

(5) The precursors obtained in (4) were heated at 350°C for 5 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 7: Preparation of Up-Converting Fluoride Phosphor with DTPA Chelator

(1) Stock solutions of GdCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 16 mL: 3.9 mL: 0.02 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 80:19.5:0.5$.

(2) 20 mL of 0.2 M diethylenetriaminepentaacetic acid (DTPA) solution was introduced to the solution prepared in (1).

(3) A solution of NaF was prepared by dissolving 2.1 g of NaF in 60 mL of de-ionized water followed by addition of 20 mL of 0.2 M EDTA solution. The solution prepared in (2) was injected into the NaF solution. The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried.

(5) The precursors obtained in (4) were annealed heated at 380°C or 8 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 8: Preparation of Up-Converting Fluoride Phosphors with HEDTA Chelator

(1) Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 15.4 mL: 4 mL: 0.6 mL and added into a 100 mL-three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 77:20:3$.

(2) 20 mL of 0.2 M hydroxyethylethylenediaminetriacetic acid (HEDTA) solution was introduced to the solution prepared in (1).

(3) A solution of NaF was prepared by dissolving 2.1 g of NaF in 75 mL of deionized water. The solution prepared in (2) was injected into the NaF solution at 100°C . The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried. The average diameter of the precursor particles was 60 nm.

(5) The precursors obtained in (4) were heated at 450°C or 1 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 9: Preparation of Up-Converting Fluoride Phosphors with DCTA Chelator

(1) Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 15.4 mL: 4 mL: 0.6 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 77:20:3$.

(2) 20 mL of 0.2 M 1,2-diaminocyclohexanetetraacetic acid (DCTA) solution was introduced to the solution prepared in (1).

(3) A solution of NaF was prepared by dissolving 2.1 g of NaF in 60 mL of deionized water. The solution prepared in (2) was injected into the NaF solution at 100°C. The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried. The average diameter of the precursor particles was 60 nm.

(5) The precursors obtained in (4) were heated at 450°C or 1 h. The nanometer-sized up-converting fluorescent particles were then obtained.

10 Example 10: Preparation of Up-Converting Fluoride Phosphors with EGTA Chelator

(1) Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 15.4 mL: 4 mL: 0.6 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 77:20:3$.

(2) 20 mL of 0.2 M ethylene glycol bis (b-aminoethylether) tetraacetic acid (EGTA) solution was introduced to the solution prepared in (1).

(3) A solution of KF was prepared by dissolving 2.1 g of KF in 60 mL of de-ionized water. The solution prepared in (2) was injected into the KF solution at 100°C. The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with de-ionized water, and dried. The average diameter of the precursor particles was 60 nm.

(5) The precursors obtained in (4) were heated at 410°C for 1 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 11: Preparation of Up-Converting Fluoride Phosphors with TTHA Chelator

(II)

25 (1) Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 15.4 mL: 4 mL: 0.6 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 77:20:3$.

(2) 20 mL of 0.2 M TTHA solution was introduced to the solution prepared in (1).

(3) A solution of NH_4F was prepared by dissolving 2.1 g of NH_4F in 60 mL of deionized water. The solution prepared in (2) was injected into the NH_4F solution at 100°C . The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with deionized water, and dried. The average diameter of the precursor particles was 60 nm.

(5) The precursors obtained in (4) were heated at 410°C for 1 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 12: Preparation of Up-Converting Fluoride Phosphors with DTPA Chelator

(1) Stock solutions of YCl_3 , YbCl_3 and HoCl_3 were mixed at a volume ratio of 14 mL: 6 mL: 0.0002 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 77:20:3$.

(2) 20 mL of 0.2 M DTPA solution was introduced to the solution prepared in (1).

(3) A solution of NaF was prepared by dissolving 2.1 g of NaF in 60 mL of deionized water. The solution prepared in (2) was injected into the NaF solution at 80°C . The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with deionized water, and dried. The average diameter of the precursor particles was 50 nm.

(5) The precursors obtained in (4) were heated at 360°C for 10 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 13: Preparation of Up-Converting Fluoride Phosphors with HEDTA Chelator (II)

(1) Stock solutions of YCl_3 , YbCl_3 and TmCl_3 were mixed at a volume ratio of 14 mL: 5.96 mL: 0.04 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Tm}^{3+} = 70:29.8:0.2$.

(2) 20 mL of 0.2 M HEDTA solution was introduced to the solution prepared in (1).

(3) A solution of NaF was prepared by dissolving 2.1 g of NaF in 60 mL of deionized water. The solution prepared in (2) was injected into the NaF solution. The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with deionized water, and dried. The average diameter of the precursor particles was 60 nm.

(5) The precursors obtained in (4) were heated at 400°C for 3 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 14: Preparation of Up-Converting Fluoride Phosphors with DCTA Chelator

(1) Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 14 mL: 6 mL: 0.001 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 70:30:0.005$.

(2) 20 mL of 0.2 M DCTA solution was introduced to the solution prepared in (1).

(3) A solution of NH_4F was prepared by dissolving 2.1 g of NH_4F in 60 mL of deionized water. The solution prepared in (2) was injected into the NH_4F solution at 80°C.

10 The mixture was stirred vigorously for 1 h at room temperature.

(4) The obtained precipitates were centrifuged, washed three times with deionized water, and dried.

(5) The precursors obtained in (4) were heated at 400°C for 4 h. The nanometer-sized up-converting fluorescent particles were then obtained.

15 Example 15: Preparation of Up-Converting Fluoride Phosphors with EGTA Chelator

(II)

(1) Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 17.4 mL: 2 mL: 0.6 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 87:10:3$.

20 (2) 20 mL of 0.2 M EGTA solution was introduced to the solution prepared in (1).

(3) A solution of NaF was prepared by dissolving 2.1 g of NaF in 50 mL of deionized water followed by addition of 0.2 M EGTA solution 10 mL. The solution prepared in (2) was injected into the NaF solution at 40°C. The mixture was stirred vigorously for 1 h at room temperature.

25 (4) The obtained precipitates were centrifuged, washed three times with deionized water, and dried.

(5) The precursors obtained in (4) were heated at 400°C for 3 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 16: Preparation of Up-Converting Fluoride Phosphors with TTHA Chelator

30 (II)

(1) Stock solutions of YCl_3 , YbCl_3 and ErCl_3 were mixed at a volume ratio of 15 mL: 2 mL: 3 mL and added into a 100 mL three-neck Erlenmeyer flask. The molar ratio of lanthanide ions in the mixture was $\text{Y}^{3+}:\text{Yb}^{3+}:\text{Er}^{3+} = 75: 10: 15$.

5 (2) A quantity of 20 mL of 0.2 M TTHA solution was introduced to the solution prepared in (1).

(3) A solution of NH_4F was prepared by dissolving 2.1 g of NH_4F in 60 mL of deionized water. The solution prepared in (2) was injected into the NH_4F solution at room temperature. The mixture was stirred vigorously for 1 h at room temperature.

10 (4) The obtained precipitates were centrifuged, washed three times with deionized water, and dried.

(5) The precursors obtained in (4) were heated at 400°C for 2 h. The nanometer-sized up-converting fluorescent particles were then obtained.

Example 17: Surface Modification of Nanometer-Scale Up-Converting Phosphors

15 An aliquot of 10mL isopropanol containing 20 mg of nanometer-scaled fluoride phosphor particles was sonicated thoroughly, until a homogenous colloid-like state of the suspension was reached. Small amount of large-sized particles settled at the bottom, and were discarded.

20 Approximately 74 mL of isopropanol was added to a 100 mL Erlenmeyer flask. It was magnetically stirred and heated to 40°C in an oil bath. The colloid-like homogenous up-converting phosphor suspension was then added into the flask, with subsequent addition of 6.75 mL of deionized water and 8.25 mL of 25% ammonium hydroxide. The flask was then sealed. After the mixture was magnetically stirred for 10 min, 0.08 mmol of tetraethyl orthosilicate (TEOS) was added into the flask and the reaction was kept on for 1 h. A quantity of 0.07 mmol of 3-aminopropyltriethoxysilane (APTES) was then added into the
25 reaction system and the suspension was stirred for an additional 3 h at 40°C . The mixture was transferred to centrifuge tubes and centrifuged at 12,000 rpm. The supernatant was discarded and pellets washed twice with deionized water. The resulting white powder was dried in oven at 60°C .

30 The APTES-modified dry powder (4mg) was re-suspended in 2 mL of 1x phosphate buffered saline (PBS buffer, 4 g sodium chloride, 0.1 g potassium chloride, 1.56 g sodium

hydrogenphosphate, 0.1 g potassium dihydrogenphosphate in 500 mL of MilliQ H₂O, pH 7.4). After a sonication period of 30 min, the suspension was added with 50 µl of 50% glutaraldehyde aqueous solution and 15 mg of sodium cyanoborohydride (NaCNBH₃) to introduce aldehyde groups onto the silanized particle surface. The reaction was kept on for 2 h at 4°C with sonication. The suspension was centrifuged and the pellet washed using 1xPBS buffer three times. The pellet was then re-suspended in 1 mL of 1x PBS buffer (CHO-modified up-converting phosphor particle stock, approximately 4 mg/mL).

Example 18: Covalent Conjugation of NeutrAvidin with Up-converting Phosphor

NeutrAvidin (2mg, Pierce) was dissolved in 0.1 mL of MilliQ H₂O. The volume was adjusted to 250 mL by addition of 1x PBS buffer.

A suspension of 0.25 mL of the 4 mg/mL CHO-modified up-converting phosphor particle stock prepared in Example 17 was sonicated for 10 min followed by the addition of the 2 mg/mL NeutrAvidin solution. Adjust the final volume to 1 mL using 1x PBS buffer. The reaction was kept on for 1 h at room temperature on a vibration station to complete the covalent conjugation of the particle to the protein molecule via aldehyde-amino group reaction. An aliquot of 5 µL of 5 M NaCNBH₃ solution was added to the mixture to reduce the Schiff base product resulted from the aldehyde-amino group reaction. After 1 h of incubation, 20 µL of 3 M tris(hydroxymethyl)aminomethane (Tris) solution was added into the system to block the excess aldehyde groups on the particle surface.

The resulting suspension was centrifuged and washed using 1x PBS buffer three times. The pellet was re-suspended into 1xPBS buffer and ready for the subsequent usage in bio-molecular assays

The above examples are included for illustrative purposes only and are not intended to limit the scope of the invention. Many variations to those described above are possible. Since modifications and variations to the examples described above will be apparent to those of skill in this art, it is intended that this invention be limited only by the scope of the appended claims.

What is claimed is:

1. A phosphor fluoride particle that emits light in the visible wavelength range when excited by long wavelength light that has a uniform particle size of less than 350 nanometers.
5. 2. The phosphor fluoride particle of claim 1, comprising a phosphor host and an absorber-emitter pair.
3. The phosphor fluoride particle of claim 2, wherein the phosphor host is selected from the group consisting of yttrium, lanthanum and gadolinium.
4. The phosphor fluoride particle of claim 2, wherein the absorber is ytterbium
10 and the emitter is selected from the group consisting of erbium, holmium, terbium and thulium.
5. The phosphor fluoride particle of claim 1, wherein the particle has a molar ratio:
(yttrium, lanthanum or gadolinium):ytterbium:(erbium, holmium, terbium or thulium)
15 = (70-90):(0-29):(0.001-15).
6. The phosphor fluoride particle of claim 1, which has a formula of $YF_3:Yb,Er$.
7. The phosphor fluoride particle of claim 1, which has a formula of $NaYF_4:Yb,Er$.
8. The phosphor fluoride particle of claim 1, which has a particle size ranging
20 from about 35 nanometers to about 200 nanometers.
9. The phosphor fluoride particle of claim 1, further comprising a transparent coating layer.
10. The phosphor fluoride particle of claim 9, wherein the transparent coating layer is SiO_2 .
- 25 11. The phosphor fluoride particle of claim 9, wherein the coated particle further comprises a layer of immobilized biological moiety.
12. A process of preparing a phosphor fluoride particle that emits light in the visible wavelength range when excited by long wavelength light that has a uniform particle size of less than 350 nanometers, which process comprises:

a) preparing an aqueous solution of soluble salts of a phosphor host, an absorber/emitter pair and a rare-earth metal chelator; and

b) contacting said prepared aqueous solution of soluble salts of said phosphor host, absorber/emitter pair and rare-earth metal chelator with an aqueous fluoride-containing compound at a temperature ranging from about 0°C to about 100°C for a sufficient time to obtain a precipitate of a phosphor fluoride particle; and

c) heating said precipitate at a temperature ranging from about 300°C to about 450°C for a time ranging from about 1 hour to about 10 hours to obtain a phosphor fluoride particle that emits light in the visible wavelength range when excited by long wavelength light that has a uniform particle size of less than 350 nanometers.

13. The process of claim 12, wherein the phosphor host is selected from the group consisting of yttrium, lanthanum and gadolinium.

14. The process of claim 12, wherein the absorber is ytterbium and the emitter is selected from the group consisting of erbium, holmium, terbium and thulium.

15. The process of claim 12, wherein the rare-earth metal chelator is selected from the group consisting of ethylenediaminetetraacetic acid, triethylenetetraaminhexaacetic acid, diethylenetriaminepentaacetic acid, hydroxyethylethylenediaminetriacetic acid, 1,2-diaminocyclohexanetetraacetic acid, ethylene glycol bis (b-aminoethylether) tetraacetic acid and a salt thereof.

16. The process of claim 12, wherein the aqueous fluoride-containing compound is selected from the group consisting of NaF, KF, NH₄F and HF.

17. The process of claim 12, wherein the aqueous fluoride-containing compound is contained in an aqueous solution prior to or concurrently with contacting with the prepared aqueous solution of soluble salts of the phosphor host, the absorber/emitter pair and the rare-earth metal chelator.

18. The process of claim 12, wherein the soluble salts of the phosphor host and the absorber/emitter pair are obtained by dissolving the corresponding metal oxide in hydrochloric acid or nitric acid and subsequently removing the residual acid.

19. The process of claim 12, wherein the amount of the rare-earth metal chelator is about 0-1 times the amount of total rare-earth ions in the aqueous solution.

20. The process of claim 12, further comprising coating the prepared phosphor fluoride particle with a transparent layer.

21. The process of claim 20, wherein the transparent layer is SiO_2 .

22. The process of claim 20, further comprising coating the transparent layer coated
5 phosphor fluoride particle with a layer of immobilized biological moiety.

23. A phosphor fluoride particle that is prepared by the process of claim 12.

24. The phosphor fluoride particle of claim 23, which has a molar ratio:
(yttrium, lanthanum or gadolinium):ytterbium:(erbium, holmium, terbium or thulium)
= (70-90):(0-29):(0.001-15).

10 25. The phosphor fluoride particle of claim 23, which has a formula of $\text{YF}_3\text{:Yb,Er}$.

26. The phosphor fluoride particle of claim 23, which has a formula of
 $\text{NaYF}_4\text{:Yb,Er}$.

27. The phosphor fluoride particle of claim 23, which has a particle size from
about 35 nanometers to about 200 nanometers.

15

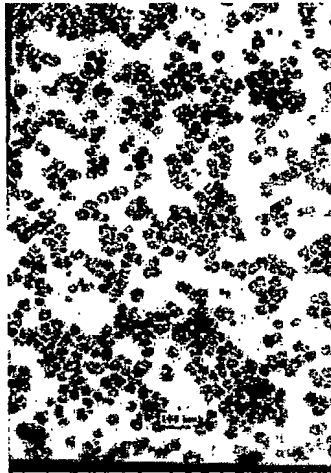


Figure 1

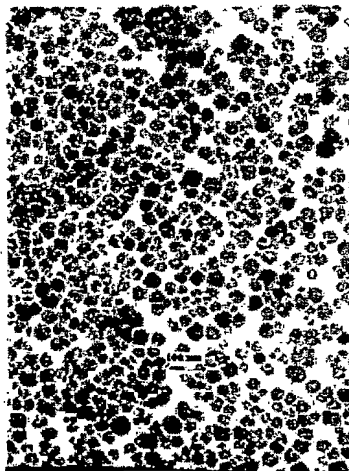


Figure 2

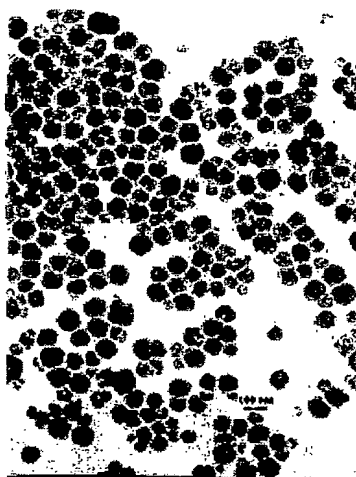


Figure 3

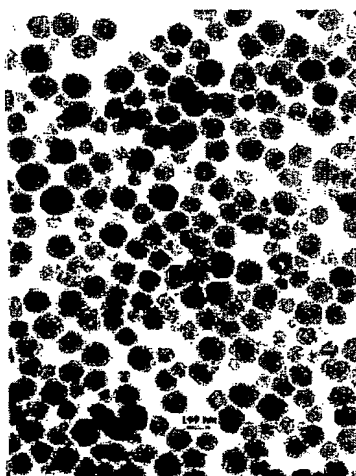


Figure 4

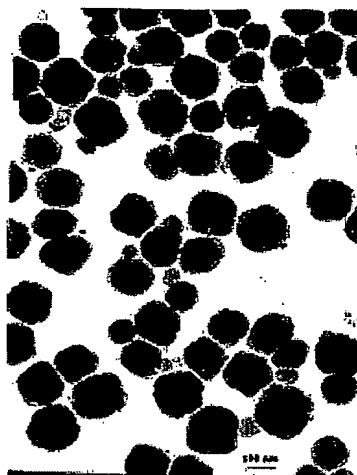


Figure 5

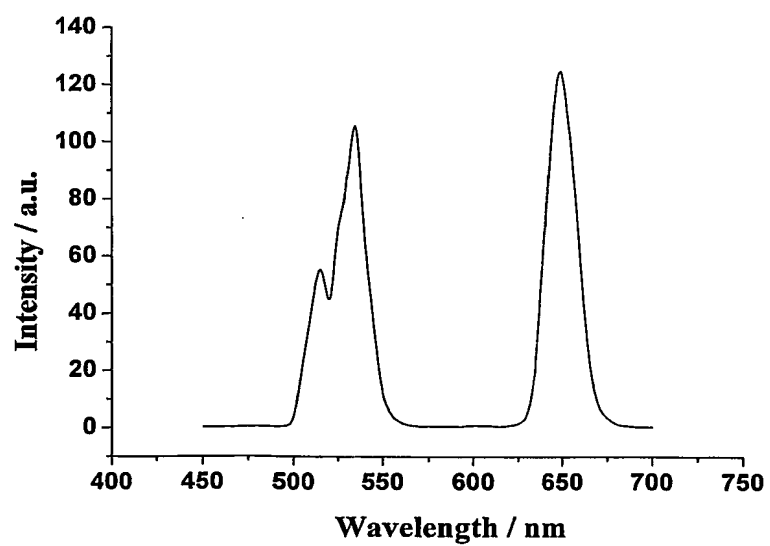


Figure 6

INTERNATIONAL SEARCH REPORT

International application No.
PCT/CN03/00268

A. CLASSIFICATION OF SUBJECT MATTER

IPC⁷: C09K11/77, C09K11/61

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC⁷: C09K11/00-11/89

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

WPI, CNPAT

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X,	CN1185475A(CHANGCHUN PHYSICS INST CHINESE ACAD SCI) 24.06.1998, Page 1, Claim 1	1-4,6-8
A		5, 9-27
P,X	CN1357599A(KERUN OPTOELECTRONIC MATERIAL SCI & TECH) 10.07.2002, Page,1,2 Example 1,4	1-4,6-8
A	JP4328191A(CENTRAL GLASS CO LTD) 17.11.92, Abstract	1-11,23-27

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

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Date of the actual completion of the international search
13.July.2003 (13.07.03)

Date of mailing of the international search report

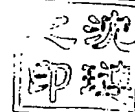
24 JUL 2003 (24.07.03)

Name and mailing address of the ISA/CN
6 Xitucheng Rd., Jimen Bridge, Haidian District,
100088 Beijing, China
Facsimile No. 86-10-62019451

Authorized officer

SHEN, Lian

Telephone No. 86-10-62093047



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